Abramsky L., Chapple J.. (1994), Prenatal diagnosis: The human side. London UK. Chapman & Hall. Book Darr A. 134-149.

Adam M.P., Fechner P.Y., Ramsdell L.A., Badaru A., Grady R.E.. (2012), Ambiguous genitalia: What Prenatal genetic testing is practicle ?. Am J med Genet A. 1337-1343.

Adiga S.K., Kalthur G., Kumar P., Girisha K.M.. (2010), Preimplantation diagnosis of genetic disease. India. J Postgrad med. 317-320.

Broke D. J.H., Rodeck C. H., Ferguson-Smith. (1992), Prenatal diagnostic and screening. New York: Churchill Livingstone. 3 - 99,741.

Crandall B.F., Golbus M.S., Goldberg J.D., Matsumoto m.. (1991), First trimester maternal serum unconjugated oestriol and alpha fetoprotein in fetal Down's syndrome. Los Angeles. Prenat. Diagn. 377-380.

Crissman H.P., Warner L., Gardner M., Schast A., Quittner A., Kogan B.. (2011), Children with disorder of sex development: A qualitative study of early prenatal experience. Int J Pediatr Endocrinol. Volume 1. 10-13.

Dewald G.W., Superbeck K.J.. (1983), Sex chromosome anomalies associated with premature gonadal failure. Semin Reprod Endocrinol. Volume 1., 79-92.

Felix M.. (1995), ISCN: An International System for Human Cytogenetic Nomenclature. 1st ed. New York. Karger. 46, 50-73.

Gardner M. R.J., Sutherland G. R.. (2004), Chromosome abnormalities and genetic counseling. 3rd ed.. 363-392.

Gogate S. (2006), Preventive Genetics. 1st ed.. Jaypee publication. 3,101,271.

Goldstein H., Nielsen K.G.. (1998), Rates and survival of individuals with trisomy 13 and 18. Clin Genet. 366- 372.

Kaur A., Singh J.R.. (2010), Chromosomal abnormalities: Genetic disease burden in India, Int J Hum Genet. 1-3.

Krishna U., Dafftary S.. (1993), Pregnancy at risk current concepts. The federation of Obst. and Gynoc. society of India (FOGSI). 157 -166.

Margaret T., Roderick R., Huntington F.. (1991), Patterns of single gene inheritance. Genetics in medicine. Volume 4. 92-97.

Milunsky A. (1992), Prenatal diagnosis of neural tube defects. In: Genetic disorder and the fetus. Plenum. New York. 379-430.

Myra C.B., Van Z., Dick L.W., Liesbeth L.L.. (2005), How unexpected are unexpected finding in prenatal cytogenetic diagnsosi ? A literature review. European Journal of Obstetrics and Gynecology and Reproductive Biology. 15-21

Nussbaum R.L., McInnes R.R., Willard H.F.. (2001), Thompson and Thompson: Genetics in Medicine. 359-391.

Nyberg D., Mahony B. S., Pretorius D. H.. (1990), Diagnostic ultrasound of fetal anomalies. Text and Atlas. Mosby: St. Louis. 38-66.

Nyhan W. L.. (1988), Diagnostic Recognition of Genetic Disease. Ann Intern Med. 503-504.

Phadke S.R., Sharda S., (2007), Uptake of invasive prenatal diagnosis tests in women after detection of soft markers for chromosomal abnormality on ultrasonographic avaluation. Indai. J Perinatol. 550-555.

Purandarey H. (2009), Essentials of Human Genetics. 2nd ed., Jaypee Brothers 327.

Purandarey H., Chakravarty A.. (2000), Human Cytogenetics Techniques And Clinical Application. 1st ed. Bhalani Publishing House. 85 – 117.

Rao Kamini A. (1996), An Introduction to genetics and fetal Medicine.. The federation of Obst. And Gynoc. society of India (FOGSI). 51-73.

Sadler T.W., Leland J. (2006), Langman's Medical Embryology. Journal of cell Biology.10th ed. Lippincott Wiliams & Wilkins. 31, 229.

Shaffer L. G., Slovak M. L., Campbell L. (2009), ISCN: An International system for human cytogenetic nomenclature. J Histochem Cytochem. 991-993.

Tan S.Y., Chan W.B., Cheng W.C., Hagarty A., Quaife K.T., (2000), Rapid prenatal diagnosis of chromosome abnormalities., Singapur Med, Volume 4. 493-497.

Thasnevis V. (1996), Prenatal diagnostic technology shaking the basics of ethics. 1stedition. Katholieke university Leuven. 76-77.

Thompson M.W., McInnes R.R., Willard H.F. (1991), Genetics in medicine. 5th edition. Philadelphia. WB Saunders. 45-84.

Verma I. C. (2001), Burden of genetic disorder in India. Indian J Pediatr. 893-898.

Verma I.C., Saxena R., Lal M., Bijarnia S., Sharma R. (2003), Genetic counseling and prenatal diagnsosis in India. Indian J Pediatr. 293-297.

Verma R.S., Babu A. (1995), Human Chromosomes: principles and techniques. 2nd ed.. New York: McGraw Hill. 11-113.

Verma R.S., Dosik H.. (1982), Recent advanced in detecting human chromosomal abnormalities by various banding techniques. Pathology Annual. Vol-17. 261-686.

Warburton D., Byrne J., Canki N. (1991), Chromosome anomalies and prenatal development: An atlas. 18-21, 57-63.

Wieacker P., Steinhard J.. (2010), The prenatal diagnosis of gentic diseas. Dtch Arztebl Int.. 857-862.

Williams J.. (1994), Amniotic cell culture and harvest. The Cytogentics Symposia. 1-8.